Billing Code 4410-09-M

## DEPARTMENT OF JUSTICE DRUG ENFORCEMENT ADMINISTRATION MANUFACTURER OF CONTROLLED SUBSTANCES NOTICE OF REGISTRATION JOHNSON MATTHEY, INC., PHARMACEUTICALS MATERIALS

By Notice dated January 6, 2012, and published in the Federal Register on January 17, 2012, 77 FR 2324, Johnson Matthey, Inc., Pharmaceuticals Materials, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydrocodone (9193)	II

Drug Schedule

Morphine (9300) II

Thebaine (9333) II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers. The Thebaine (9333) will also be used to manufacture other controlled substances in bulk which will also be for sale in bulk to its customers.

No comments or objections have been received. DEA has considered the factors in 21 USC § 823(a) and determined that the registration of Johnson Matthey Inc.,

Pharmaceuticals Materials to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Johnson Matthey Inc., Pharmaceuticals Materials to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and

local laws, and a review of the company's background and history.

Therefore, pursuant to 21 USC § 823(a), and in accordance with 21 CFR § 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Joseph T. Rannazzisi
Deputy Assistant Administrator
Office of Diversion Control
Drug Enforcement Administration

DATED: May 11, 2012

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